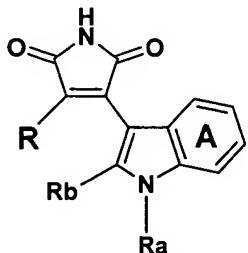


Listing of Claims

This listing of claims will replace all prior versions, and listings, of claims in the application:

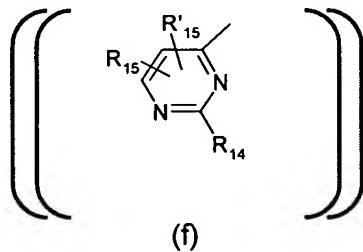
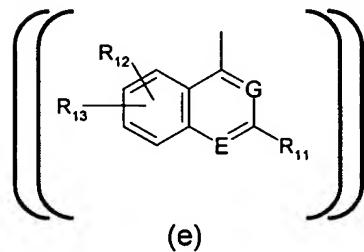
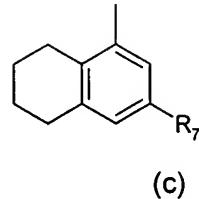
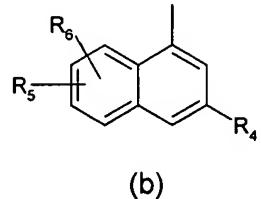
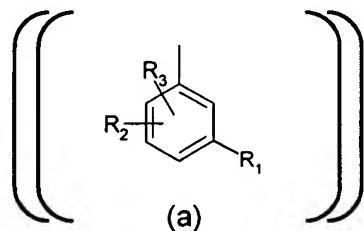
1. (currently amended) A compound of formula I



wherein

R_a is H; C₁₋₄alkyl; or C₁₋₄alkyl substituted by OH, NH₂, NHC₁₋₄alkyl or N(C₁₋₄ alkyl)₂; R_b is H; or C₁₋₄alkyl;

R is a radical of formula (a) [[,]] (b) [[,]] or (c), (e) or (f)



wherein

each of R_4 , $R_{4[[\cdot]]}$ and $R_{7[[\cdot]]}$ R_{14} and R_{14} is OH; SH; a heterocyclic residue; $NR_{16}R_{17}$

wherein each of R_{16} and R_{17} , independently, is H or C_{1-4} alkyl or R_{46} and R_{47} form together with the nitrogen atom to which they are bound a heterocyclic residue; or a radical of formula □



wherein X is a direct bond, O, S or NR_{18} wherein R_{18} is H or C_{1-4} alkyl, R_c is C_{1-4} alkylene or C_{1-4} alkylene wherein one CH_2 is replaced by CR_xR_y wherein one of R_x and R_y is H and the other is CH_3 , each of R_x and R_y is CH_3 or R_x and R_y form together $-CH_2-CH_2-$, and

Y is bound to the terminal carbon atom and is selected from $OH[[\cdot]]$ a heterocyclic residue and $-NR_{19}R_{20}$ wherein each of R_{19} and R_{20} independently is H, C_{3-6} cycloalkyl, C_{3-6} cycloalkyl- C_{1-4} alkyl, aryl- C_{1-4} alkyl or C_{1-4} alkyl optionally substituted on the terminal carbon atom by OH, or R_{49} and R_{20} form together with the nitrogen atom to which they are bound a heterocyclic residue;

each of R_2 , $R_3[[\cdot]]$ $R_5[[\cdot]]$ and $R_6[[\cdot]]$ R_{42} , R_{43} , R_{45} and R'_{45} , independently, is H, halogen, C_{1-4} alkyl, CF_3 , OH, SH, NH_2 , C_{1-4} alkoxy, C_{1-4} alkylthio, NHC_{1-4} alkyl, $N(C_{1-4}$ alkyl) $_2$ or CN;

E is $N=$ and G is $CH=$; and

ring A is optionally substituted,
or a salt thereof.

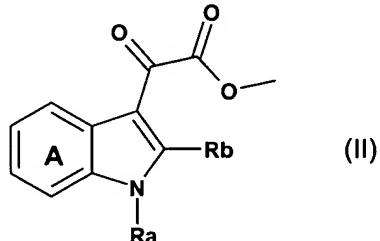
2. - 4. (cancelled)

5. (withdrawn) A compound according to claim 1 wherein R is a radical of formula (e) or (f).

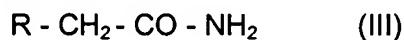
6. (canceled)

7. (withdrawn-currently amended) A process for the preparation of a compound of formula I according to claim 1 which process comprises

- a) reacting a compound of formula II

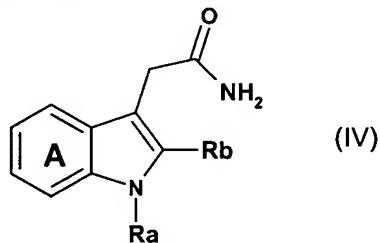


wherein R_a, R_b and ring A are as defined in claim 1,
with a compound of formula III



wherein R is as defined in claim 1,

- b) reacting a compound of formula IV



wherein R_a, R_b and ring A are as defined in claim 1,
with a compound of formula V



wherein R is as defined in claim 1; or

- c) converting in a compound of formula I a substituent R₄, R₄_[1,1] or R₇_[1,1] R₈, R₁₁ or R₁₄
into another substituent R₄, R₄_[1,1] or R₇_[1,1] R₈, R₁₁ or R₁₄

and, where required, converting the resulting compound of formula I obtained in free form to a salt form or vice versa, as appropriate.

8. (canceled)

9. (original) A pharmaceutical composition comprising a compound of formula I according to claim 1 in free form or pharmaceutically acceptable salt form in association with a pharmaceutically acceptable diluent or carrier therefor.

10. (canceled)

11. (withdrawn) A method for preventing or treating disorders or diseases mediated by T lymphocytes and/or PKC in a subject in need of such treatment, which method comprises administering to said subject an effective amount of a compound of formula I according to claim 1 or a pharmaceutically acceptable salt thereof.